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<p>(54) Title: PHARMACEUTICAL DIPEPTIDE COMPOSITIONS AND METHODS OF USE THEREOF</p>			
<div style="text-align: center;"> <p>(I)</p> </div>			
<p>(57) Abstract</p> <p>Methods are provided for the therapy of immunodeficient, immunodepressed or hyperactive immune states and for the prevention and treatment of opportunistic infections in such states comprising administering to a subject a pharmaceutically acceptable composition comprising as an active ingredient the dipeptide L-Glu-L-Trp, the cyclic monomer thereof, polymers thereof of the formula <math>H_2N - [L-Glu-L-Trp]_n - CO_2H</math>; cyclic polymers thereof of formula (I) and their pharmaceutically acceptable salts thereof, wherein n and m are independently <math>\geq 2</math>.</p>			

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**PHARMACEUTICAL DIPEPTIDE COMPOSITIONS  
AND METHODS OF USE THEREOF**

The present invention is directed to dipeptide pharmaceutical compositions and uses thereof, in particular, uses thereof for treatment of immunodepressed states and of opportunistic infections in immunodepressed states.

BACKGROUND OF THE INVENTION

Several polypeptides found in the thymus gland have been implicated as playing roles in the development and maintenance of immunological competence in animals, including human beings. Some of these polypeptides have been shown to stimulate the maturation, differentiation and function of T-cells. For example, a heat-stable fraction isolated from calf thymus extracts, designated as Thymosin fraction 5, has been shown to reconstitute immune functions in thymic-deprived or immunodepressed individuals. Several peptides have been isolated from Thymosin fraction 5, such as Thymosin  $\alpha_1$  (28 amino acids, U.S. Patent No. 4,079,127), Thymosin  $\beta_4$  (44 amino acids, Low et al., PNAS, 78,1162-1166 (1981)), Thymosin  $\beta_8$  (39 amino acids, U.S. Patent No. 4,389,343) and Thymosin  $\beta_9$  (41 amino acids, U.S. Patent No. 4,389,343). However, practical administration of such polypeptides is expensive due

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to the relatively low yield and complexity of isolation and/or manufacture of such long chain polypeptides. Most importantly, in some cases, these polypeptides produce side reactions in patients.

5 The present invention is based in part on the discovery that a dipeptide, hereinafter referred to as Thymogen, exhibits a broad range of efficacy for prevention and treatment of opportunistic infections in immunodepressed states, and for therapeutically  
10 effective treatment of immunodeficient states. This is believed to be highly unexpected for such a relatively small compound to exhibit such a broad range of activity. Furthermore, we have not found any significant side effects from the use of the  
15 dipeptide according to the present invention. Due to its simple nature, the dipeptide is rather inexpensive to manufacture.

As used herein, the terms "immunomodulator" and "immunomodulating" encompass the activity of  
20 enhancing or restoring the subject's immune system, as evidenced by measurable blood parameters and/or the patient's improved ability to combat infection or disease, and the ability to heal tissue. Hence, immunomodulation encompasses improvement of the  
25 immune system due to an immunodeficient state (for example, caused by removal of the thymus), and/or an immunodepressed state (for example, caused by exposure to radiation). Furthermore, the present invention provides for modulation of the immune  
30 system by lowering blood parameters and other indicia of the immune state if these indicia are abnormally elevated. The present invention encompasses the therapeutic method of treating the immunodeficient, immunodepressed or elevated immune state per se, thus  
35 providing prophylaxis against infection and disease,

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as well as a treatment of infection, disease or wound indirectly by enhancing the immune system.

It is therefore an object of the present invention to provide pharmaceutical compositions of the dipeptide Thymogen which have broad immunomodulating activity, as well as activity for other uses such as treatment of infections, disease and wounds (burns, frost bites, and the like), enhancement of metabolic processes, and many other uses.

10 It is an object of the present invention to provide therapeutic methods for treatment of immunodepressed and immunodeficient states.

It is yet another object of the present invention to provide methods for preventing and treating  
15 opportunistic infections in immunodeficient and immunodepressed states.

These and other objects will be apparent from the following description and appended claims.

#### DESCRIPTION OF THE PREFERRED EMBODIMENTS

20 The present invention provides pharmaceutical preparations comprising the dipeptide L-Glu-L-Trp, including cyclic forms and linear and cyclic polymers of the dipeptide, using the normal convention wherein the first named amino acid is the amino terminus and  
25 the last named amino acid is the carboxyl terminus. For convenience, all forms will be collectively referred to herein as "the dipeptide". The compositions containing the dipeptide according to the present invention may be formulated into any  
30 convenient formulation which allows for the active ingredient to be absorbed into the blood stream.

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Intramuscular and intranasal forms of application are preferred. The preferred dosage rate of the active ingredient for intramuscular administration is about 50 to 100 $\mu$ g per dose for adults (for a 300 to 1000 $\mu$ g total treatment therapy); for infants up to 1 year old about 10 $\mu$ g per dose, for infants 1 to 3 years old about 10 to 20 $\mu$ g per dose; for infants 4 to 6 years old about 20 to 30 $\mu$ g per dose, for children 7 to 14 years old about 50 $\mu$ g per dose. All of the foregoing dosages are useful for a treatment of 3 to 10 days, depending upon the immunodeficiency level. The treatment may be repeated as needed, usually within 1 to 6 months.

For prophylactic uses against opportunistic infections in immunodeficient or immunodepressed patients, the intramuscular and/or intranasal single daily dose for adults may be from about 50 to 10 $\mu$ g, and for children about 10 to 50  $\mu$ g per dose for treatment over 3 to 5 days.

For treatment of burns, frost bite, or other wounds, including chronic apical periodontitis, the dipeptide may be applied in about 100 $\mu$ g doses as a paste or other suitable medium.

For ophthalmology, such as for treatment of infectious eye diseases, the dipeptide may be applied in single daily dosages of about 10 $\mu$ g (over 4 to 10 days) or as installations into the conjunctival cavity at about 5 $\mu$ g twice daily over about 4 to 5 days.

The dipeptide may be utilized intramuscularly as an injection solution with the active ingredient in a therapeutically effective immunopotentiating amount of about .001 to .01% by weight. If presented in the

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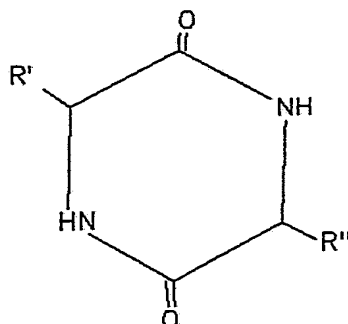
form of a tablet, capsule or suppository it is preferred that the active ingredient be present in an amount of about 0.1mg per tablet, suppository or capsule. If presented in such form, the capsule, 5 suppository or tablet may also contain other conventional excipients and vehicles such as fillers, starch, glucose, etc.

The dipeptide may be obtained by conventional peptide synthesis, including the Merrifield solid 10 state peptide synthesis technique. Typically an amino and side chain protected derivative of an activated ester of glutamic acid is reacted with protected L-tryptophan. After elimination of the protecting groups and conventional purification, such 15 as by thin layer or GL chromatography, the peptide may be purified such as by, lyophilization, gel purification, and the like.

The purified dipeptide L-Glu-L-Trp, comprises a white powder (if lyophilized; otherwise, it is 20 crystalline), soluble in water, DMF; insoluble in chloroform and ether.  $[\alpha]_D^{22} = +12.6$ ;  $C = 0.5$  H<sub>2</sub>O.  $R_f = 0.65$  (butanol: acetic acid: water = 3:1:1). UV ( $275 \pm 5$ nm, max). NMR (500MHz): 0.001mol/l of the peptide solution, Trp (3.17; 3.37; 25 4.57; 7.16; 7.24; 7.71; 7.49); Glu (1.90; 1.96; 2.21; 3.72).

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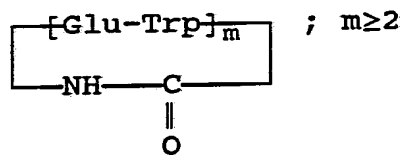
Other forms of the dipeptide are also encompassed by the invention, including the cyclized monomer.



Wherein R' and R'' are, respectively, the alpha-side chains of Glu and Trp;

5 Linear polymers  $\text{H}_2\text{N} - [\text{Glu-Trp}]_n - \text{CO}_2\text{H}$ ,  $n \geq 2$

and cyclic polymers



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all of these forms, including the dipeptide monomer, will be referred to as the "dipeptide". The linear polymers may be made by conventional peptide synthesis, including Merrifield solid-state peptide methodology. The cyclic monomer and polymers may be prepared by cyclizing the linear peptide in with peptide linkage agents in dilute solutions.

The active dipeptide ingredient of the pharmaceutical preparations according to the present invention may be used as a free peptide or in the form of a water soluble pharmaceutically acceptable salt, such as a sodium, potassium, ammonium or zinc salt. It will be understood that the dipeptide may be administered with other active ingredients which independently impart an activity to the composition, such as, antibiotics, interferon, anesthetics, and the like.



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The most preferred formulation according to the present invention is a solution for intramuscular injection containing about .001 to .01% by weight (.0001-.001mg/kg body weight, or 10-100 $\mu$ g active ingredient per 1ml solvent). The pharmaceutically acceptable vehicle for this injection form may be any pharmaceutically acceptable solvent such as 0.9% aqueous sodium chloride, distilled water, Novocaine solution, Ringer's solution, glucose solution, and the like. The dipeptide containing compositions according to the present invention may be administered in a compatible pharmaceutical suitable for parenteral administration (e.g., intravenous, subcutaneous, intramuscular). The preparations may be subjected to conventional pharmaceutical operations, such as sterilization, and may contain adjuvants, such as preservatives, stabilizers, wetting agents and the like.

The pharmaceutical preparations according to the present invention demonstrate a high effectiveness in the treatment of immunodepressed and immunodeficient states for the preventing and treatment of opportunistic infections in those states.

Also included within the scope of the present invention are the pharmaceutically acceptable salts of the dipeptide, such as sodium or potassium or strong organic bases, such as guanidine.

The dipeptide containing compositions according to the present invention have activity in the restoration and stimulation of the immune functions. Thus they are useful in the treatment of opportunistic infections of an immunodepressed subject in an immunopotentiating effective amount as described above.

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The dipeptide compositions according to the present invention may also be used in veterinary practice as an immunomodulatory agent for prophylaxis and treatment of hypotrophy in farming  
5 animals, fur bearing animals and poultry.

Among the opportunistic infections which may be treated utilizing the compositions according to the present invention are: respiratory diseases, influenza, AIDS, burns, wounds, other open sores,  
10 rashes (due to allergic reactions), sun exposure, local trauma (with an ointment), eczemas, psoriasis, and the like. Furthermore, the compositions according to the present invention may be utilized to assist healing in immunodepressed or immunodeficient  
15 states, such as for the healing of bone fractures, lesions, gingival diseases, gynecological infections, infralymphatic infections, and the like. The compositions may also be used to enhance the immunodeficient state to increase susceptibility to  
20 microbial antibiotics and to enhance the patient's responsive reaction to other types of therapies.

The compositions according to the present invention also may be utilized to enhance metabolic processes; to enhance production of blood insulin;  
25 for treatment of irradiated cancer patients, as well as for veterinary uses.

Another important use is the treatment of chronic fatigue syndrome (CFS), which is believed to be a manifestation of an immunodepressed state.

30 The following examples are provided to further elucidate the invention, but are not intended to restrict the invention in scope or spirit in any way.

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EXAMPLE 1

About 262 adult patients were treated over a period of 2 months on a daily basis with intramuscular injections of solutions containing 100  $\mu$ g of Thymogen. These patients were treated for 2 months immediately succeeding exposure to radiation caused from the Chernobyl nuclear accident. As a control, about 18 people exposed to radiation were tested for various blood parameters to establish a baseline.

The results are shown below.

THE EFFICIENCY OF RADIATION IMMUNODEFICIENCY CORRECTION  
TWO MONTHS AFTER IRRADIATION  
( $\bar{X} \pm m$ )

Indices	Examined groups		
	Healthy (control)	Irradiated	
		Prior to therapy	After THYMOGEN therapy
Leukocytes, abs	5.6 $\pm$ 0.8	3.5 $\pm$ 0.4*	5.0 $\pm$ 1.2**
Lymphocytes, abs	1.98 $\pm$ 0.16	0.80 $\pm$ 0.24*	1.9 $\pm$ 0.4**
CD2-DR+, %	35.8 $\pm$ 0.9	21 $\pm$ 4*	30.0 $\pm$ 1.2**
CD2-DR+, abs	0.59 $\pm$ 0.04	0.16 $\pm$ 0.04*	0.55 $\pm$ 0.06*
CD2, %	49.3 $\pm$ 1.5	32 $\pm$ 7	48.7 $\pm$ 1.8**
CD2, abs	0.98 $\pm$ 0.09	0.55 $\pm$ 0.08*	1.13 $\pm$ 0.08**
E-RFC, %	30.2 $\pm$ 1.6	22.9 $\pm$ 1.9*	27.4 $\pm$ 2.4**
LMI with ConA, %	65.0 $\pm$ 2.1	120 $\pm$ 17*	90 $\pm$ 10**
CD19, %	22.0 $\pm$ 1.7	32 $\pm$ 3*	27 $\pm$ 4
CD19, abs	0.46 $\pm$ 0.03	0.26 $\pm$ 0.06*	0.51 $\pm$ 0.10**
IgM, g/l	1.1 $\pm$ 0.4	0.87 $\pm$ 0.07	1.00 $\pm$ 0.10
IgG, g/l	11.1 $\pm$ 0.9	10.2 $\pm$ 2.0	10.0 $\pm$ 1.0
IgA, g/l	1.70 $\pm$ 0.10	1.5 $\pm$ 0.4	1.49 $\pm$ 0.19

\* - statistically significant ( $P < 0.05$ ) vs. the indices in healthy people;

\*\* - statistically significant ( $P < 0.05$ ) vs. the data obtained prior to immunocorrection with THYMOGEN;

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abs - cell concentration presented as  $10^9/l$ ;  
 LMI - leucocyte migration inhibition;  
 RFC - rosette-forming cells.

EXAMPLE 2

5 The group of patients described in Example 1 were further treated for a period of 36 months and tested again subsequent to the first stage of therapy (after 4 months) and after the second stage of therapy (6 months). The blood parameters are shown below. As  
 10 can be seen most of the blood parameters were elevated after both the first and second stages of therapy.

THE PROLONGED THYMOGEN THERAPY TRIALS RESULTS  
 IN IRRADIATED PATIENTS  
 (X $\pm$ m)

15

Indices	Interims of examination		
	prior to therapy	after the 1st stage of THYMOGEN use	after the 2nd state of THYMOGEN use
Leucocytes, abs	3.5 $\pm$ 0.5	4.7 $\pm$ 0.2*	5.5 $\pm$ 0.3*
Lymphocytes, abs	1.0 $\pm$ 0.5	1.5 $\pm$ 0.4	1.9 $\pm$ 0.5*
CD2-DR+, %	12.8 $\pm$ 2.6	22.3 $\pm$ 0.5	29 $\pm$ 3*
20 CD2-DR+, abs	0.13 $\pm$ 0.04	0.34 $\pm$ 0.05*	0.56 $\pm$ 0.08*
CD3, %	24 $\pm$ 3	35 $\pm$ 4*	46 $\pm$ 3*
CD3, abs	0.26 $\pm$ 0.05	0.49 $\pm$ 0.06*	0.89 $\pm$ 0.11*
CD4, %	7.1 $\pm$ 1.1	19.5 $\pm$ 1.7*	24.1 $\pm$ 1.5*
CD4, abs	0.07 $\pm$ 0.01	0.28 $\pm$ 0.03*	0.45 $\pm$ 0.04*
25 CD8, %	17 $\pm$ 3	15.4 $\pm$ 2.3	22.3 $\pm$ 2.2*
CD8, abs	0.16 $\pm$ 0.04	0.23 $\pm$ 0.03	0.40 $\pm$ 0.05*
CD19, %	12.2 $\pm$ 1.9	15.0 $\pm$ 2.8	21.1 $\pm$ 2.1*
CD19, abs	0.14 $\pm$ 0.04	0.21 $\pm$ 0.06	0.39 $\pm$ 0.06*

30 \* - statistically significant (P<0.05) in comparison with the indices prior to therapy;

abs - cell concentration presented as  $10^9/l$ .

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EXAMPLE 3

The patients described in Example 1 were tested for blood parameters the first few days after exposure to the radiation of the Chernobyl accident.

- 5 It could be seen from the table below that response to the treatment was observed even after a few weeks? of treatment.

THYMOGEN INFLUENCE ON IMMUNE STATUS IN EARLY TERMS  
AFTER IRRADIATION AFFECTION  
( $\bar{X} \pm m$ )

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Indices	Examined groups		
	Healthy (control)	Irradiated	
		Prior to therapy	After THYMOGEN therapy
Leukocytes, abs	5.7 $\pm$ 0.3	3.8 $\pm$ 0.3*	6.4 $\pm$ 0.8**
Lymphocytes, abs	1.91 $\pm$ 0.12	1.15 $\pm$ 0.14*	2.27 $\pm$ 0.16**
CD2-DR+, %	30.8 $\pm$ 1.1	17.6 $\pm$ 2.0*	31 $\pm$ 3**
CD2-DR+, abs	0.59 $\pm$ 0.04	0.20 $\pm$ 0.03*	0.69 $\pm$ 0.08**
CD2, %	50.6 $\pm$ 1.6	47 $\pm$ 4	50.9 $\pm$ 2.4
CD2, abs	0.98 $\pm$ 0.09	0.55 $\pm$ 0.08*	1.13 $\pm$ 0.07**
E-RFC, %	29.7 $\pm$ 2.5	29.8 $\pm$ 2.6	23.4 $\pm$ 2.6
LMI with ConA, %	66 $\pm$ 4	98 $\pm$ 9*	60 $\pm$ 7**
CD19, %	22.8 $\pm$ 2.2	27.0 $\pm$ 2.8	30.5 $\pm$ 1.9*
CD19, abs	0.47 $\pm$ 0.03	0.30 $\pm$ 0.05*	0.68 $\pm$ 0.04**
IgM, g/l	1.1 $\pm$ 0.4	0.51 $\pm$ 0.08*	0.58 $\pm$ 0.10*
IgG, g/l	10.1 $\pm$ 0.9	8.6 $\pm$ 1.3	9.2 $\pm$ 0.7
IgA, g/l	1.71 $\pm$ 0.16	2.07 $\pm$ 0.20	1.11 $\pm$ 0.09*, **
C3, g/l	0.57 $\pm$ 0.03	0.74 $\pm$ 0.07	0.68 $\pm$ 0.04

\* - statistically significant ( $P < 0.05$ ) in comparison with the indices in healthy people

\*\* - statistically significant ( $P < 0.05$ ) in comparison with the data obtained prior to THYMOGEN use;

30 LMI - leukocyte migration inhibition;

abs - cells concentration presented as  $10^9/l$ .

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EXAMPLE 4

A number of (36) breast cancer patients were treated with the Thymogen by injection of daily dosages of 100  $\mu$ g (a.i.). The patients had been previously treated with radiation therapy (single doses 2 grad; total dose 45-50 grad). It can be seen from the table below the treatments restore their blood parameter levels.

10 IMMUNITY AND NON-SPECIFIC RESISTANCE INDICES  
IN BREAST CANCER PATIENTS  
TREATED WITH THYMOGEN AFTER RADIOTHERAPY

(X $\pm$ m)

Indices	Prior to radiotherapy	After radiotherapy	After THYMOGEN use
15 Lymphocytes ( $\times 10^9/l$ )	1.61 $\pm$ 0.18	0.79 $\pm$ 0.09*	1.72 $\pm$ 0.21**
T-lymphocytes ( $\times 10^9/l$ )	0.83 $\pm$ 0.07+	0.32 $\pm$ 0.03*	0.92 $\pm$ 0.12**
"Active" T-lymphocytes ( $\times 10^9/l$ )	0.49 $\pm$ 0.06	0.19 $\pm$ 0.03*	0.52 $\pm$ 0.07**
20 T-helpers (OKT4 <sup>+</sup> ) ( $\times 10^9/l$ )	0.30 $\pm$ 0.03	0.12 $\pm$ 0.01*	0.39 $\pm$ 0.04**
T-suppressors (OKT8 <sup>+</sup> ) ( $\times 10^9/l$ )	0.28 $\pm$ 0.04	0.16 $\pm$ 0.02*	0.21 $\pm$ 0.03
OKT4 <sup>+</sup> /OKT8 <sup>+</sup>	1.07 $\pm$ 0.09	0.75 $\pm$ 0.06*	1.86 $\pm$ 0.17**
DSH <sup>a</sup> to tuberculin (mm)	7.3 $\pm$ 0.4	2.6 $\pm$ 0.2*	8.7 $\pm$ 0.6**
25 LMI <sup>b</sup> with ConA (%)	68 $\pm$ 4	96 $\pm$ 7*	71 $\pm$ 5**
SI <sup>c</sup> to THYMOGEN	1.23 $\pm$ 0.15	1.19 $\pm$ 0.13	1.27 $\pm$ 0.14**
B-lymphocyte (Ig <sup>+</sup> ) ( $\times 10^9/l$ )	0.15 $\pm$ 0.02	0.11 $\pm$ 0.01	0.17 $\pm$ 0.02
Phagocytic index	4.3 $\pm$ 0.3	2.06 $\pm$ 0.18*	3.7 $\pm$ 0.2**
30 Cation proteins	1.58 $\pm$ 0.09	1.36 $\pm$ 0.08*	1.49 $\pm$ 0.12
C <sub>3</sub> -complement (g/l)	0.75 $\pm$ 0.05	0.66 $\pm$ 0.04	0.68 $\pm$ 0.04

- \* - statistically significant (P<0.05) vs. the analogous index before radiotherapy;  
 \*\* - statistically significant (P<0.05) vs. the analogous index after radiotherapy;  
 35 a - Delayed-Skin Hypersensitivity;  
 b - Leukocyte Migration Inhibition;  
 c - Sensitivity Index.

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EXAMPLE 5

On peripheral blood of human volunteers in vitro. Cell cultures were incubated and treated. As can be seen from the table below, after 24 hrs. incubation at concentrations of 1  $\mu\text{g/ml}$  and 100  $\mu\text{g/ml}$ , there was statistically no mutagenic effect in these cultures.

THE CALCULATION OF CHROMOSOME STRUCTURAL DAMAGES  
IN HUMAN PERIPHERAL BLOOD LYMPHOCYTES

Dose of the medicine	Number of analysed metaphases	Metaphases with chromosome structural aberrations		Index of re-liability (P)	Level of mutagenic effect (numbers)
		#	%		
Control	1000	15	1.5	-	-
THYMOGEN - 1 $\mu\text{g/ml}$	1000	15	1.5	>0.05	0
THYMOGEN - 100 $\mu\text{g/ml}$	1000	16	1.6	>0.05	0

EXAMPLE 6

About 263 patients were treated with doses of 100  $\mu\text{g}$  of Thymogen introduced intramuscularly on a daily basis over a period of 3 years after exposure to the radiation at the Chernobyl accident. Blood parameters after 3 years of such treatment were restored to the statistical norm prior to their exposure to the radiation.

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	Indices	Statistical norma	Results of victims examination		
			Prior to therapy	After therapy	
				THYMOGEN	Conventional
5	Leucocytes, abs	5.2±0.2	5.8±0.3	5.6±0.4	5.5±1.0
	Lymphocytes, abs	1.96±0.06	2.0±0.3	2.1±0.3	1.8±0.23
	CD2-DR+, %	30.8±1.1	15 ± 3 *	32 ± 3 **	18.4±2.5
	CD2-DR+, abs	0.59±0.04	0.30±0.06 *	0.66±0.10 **	0.34±0.11
	CD3, %	55.6±1.9	67.7±2.7 *	59.2±2.1 **	61 ± 3 *
10	CD3, abs	1.09±0.08	1.33±0.05 *	1.21±0.15	1.12±0.18
	CD4, %	35.3±2.7	36.7±2.6	36.2±1.7	38 ± 3
	CD4, abs	0.69±0.05	0.72±0.05	0.74±0.08	0.70±0.05
	CD8, %	21.3±0.9	29.7±0.9 *	23.2±2.1 **	25.0±2.7 **
	CD8, abs	0.41±0.03	0.56±0.02 *	0.48±0.07 **	0.46±0.06 **
15	T4/T8	1.64±0.12	1.24±0.10 *	1.58±0.04 **	1.52±0.13
	LI %	59.7±1.7	140 ± 30 *	75 ± 6 **	107 ± 10 **
	B-Ig+, %	13.8±1.2	10.7±0.3	11.0±0.3	11.2±0.7
	B-Ig+, abs	0.29±0.02	0.21±0.01	0.23±0.04	0.20±0.05
	B-IgM+, %	6.4±0.7	3.0±0.3 *	4.1±0.6 *	4.4±0.3
20	B-IgM+, abs	0.12±0.01	0.062±0.002	0.12±0.003 *	0.08±0.004
	B-IgG+, %	4.1±0.5	4.7±0.9	4.8±0.5	4.6±0.5
	B-IgG+, abs	0.078±0.008	0.059±0.003	0.09±0.007	0.08±0.006
	B-IgA+, %	2.2±0.2	2.3±0.3	1.9±0.3	1.98±0.09
	B-IgA+, abs	0.041±0.004	0.048±0.006	0.04±0.002	0.04±0.002
25	IgM, g/l	1.15±0.06	0.53±0.09 *	1.06±0.06 **	1.03±0.13 **
	IgG, g/l	11.5±0.5	13.2±1.1	10.9±1.3	11.3±1.2
	IgA, g/l	1.90±0.08	0.82±0.25 *	1.2±0.4 *	1.1±0.3

\* - statistically significant (P<0.05) vs. the indices in healthy persons;

30 \* - statistically significant (P<0.05) vs. the data obtained prior to therapy;

abs - cell concentration presented as 10<sup>9</sup>/l;

IMI - leucocyte migration inhibition.



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EXAMPLE 7

The patients described above in Example 6 were tested for blood parameters after 6 months of treatment immediately following exposure to the  
5 radiation caused by the Chernobyl accident. The results below show that after 6 months those treated with Thymogen showed improvement over those patients who were not treated.

THYMOGEN USE EFFICIENCY IN ACUTE RADIATION SICKNESS  
(6 MONTHS AFTER ACCIDENT)  
( $\bar{X} \pm m$ )

Indices	Healthy people (statistic norma)	Patients (suffered in accidence)			
		Prior to therapy	After therapy		
			Without THYMOGEN	With THYMOGEN	
5	Lymphocytes, %	33.9±1.2	32.9±2.4	29.2±2.0	30.0±1.8
	Lymphocytes, abs	1.96±0.06	1.49±0.14*	1.39±0.13	1.52±0.12*
	CD2, %	53.6±1.9	38.7±2.7*	32 ± 3*	49 ± 3**
	CD2, abs	1.05±0.05	0.56±0.04*	0.44±0.04*	0.75±0.05*, **
10	CD2-DR+, %	30.8±1.1	18.9±1.6*	19.7±1.2*	20.8±1.6*, **
	CD2-DR+, abs	0.59±0.04	0.30±0.25*	0.28±0.02*	0.31±0.02*, **
	CD3, %	55.6±1.9	39.0±2.4*	37 ± 5*	53.4±1.8**
	CD3, abs	1.09±0.08	0.58±0.04*	0.51±0.03*	0.82±0.04**
	CD4, %	35.3±2.7	20.3±1.3*	18.9±1.3*	32.6±1.4**
15	CD4, abs	0.69±0.05	0.30±0.03*	0.26±0.03*	0.50±0.04**
	CD8, %	21.3±0.9	19.5±1.5	17.5±1.6	21.2±1.8
	CD8, abs	0.41±0.03	0.29±0.03	0.24±0.03	0.32±0.03
	T4/T8	1.64±0.12	1.04±0.04*	1.08±0.10*	1.54±0.11**
	LMI	59.7±1.7	106 ± 6*	107 ± 6*	72.7±4.5*, **
20	CD19, %	25.00±0.12	18.2±2.1*	23 ± 3	26.7±2.1
	CD19, abs	0.49±0.04	0.27±0.03*	0.31±0.05*	0.41±0.03**
	B-Ig+, %	13.8±1.2	15.8±1.3	16.2±1.7	19.0±1.3*, **
	B-Ig+, abs	0.29±0.02	0.23±0.03	0.23±0.04	0.29±0.03
	B-IgM+, %	6.4±0.7	6.3±0.8	5.4±0.5	8.8±0.7
25	B-IgM+, abs	0.12±0.01	0.09±0.01*	0.08±0.01*	0.13±0.02
	B-IgG+, %	4.1±0.5	7.8±0.9*	7.1±0.8*	6.4±0.5*
	B-IgG+, abs	0.082±0.008	0.098±0.007	0.104±0.008	0.100±0.007
	B-IgA+, %	2.20±0.20	1.80±0.15*	1.70±0.20*	1.8±0.3
	B-IgA+, abs	0.038±0.004	0.033±0.004	0.024±0.003	0.030±0.002
30	IgM, g/l	1.15±0.06	1.14±0.08	1.20±0.07	1.07±0.09
	IgG, g/l	11.5±0.5	11.9±1.0	11.7±0.9	10.9±1.1
	IgA, g/l	1.9±1.0	1.6±0.8	1.6±0.8	1.8±0.9

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- \* - statistically significant ( $P < 0.05$ ) vs. the indices in healthy people;
- \*\* - statistically significant ( $P < 0.05$ ) vs. the data obtained before immunocorrection;
- 5 abs - cell concentration presented as  $10^6/l$ ;
- LMI - leucocyte migration inhibition.

#### EXAMPLE 8

A group of 452 persons were treated with daily dosages of 100  $\mu$ g of Thymogen administered  
10 intramuscularly over a period of 5-10 days and compared with a random group (250 persons) (not similarly treated) as a control. The cases of respiratory diseases and influenza were recorded for both groups. As can be seen from the table below,  
15 the untreated group had a greater occurrence of the diseases and sicknesses, hospitalization or disablement than the group treated with Thymogen.

**THYMOGEN CLINICO-EPIDEMIOLOGICAL  
PREVENTIVE EFFICIENCY IN ARD AND FLU**

	Indices	Group of observation		Index of efficiency
		THYMOGEN	Control	
5	Sickness rate per 100 persons/month	9.8	30.4	3.1
	Pneumonia rate/100 persons/month	0.20	0,50	2.5
	Need in hospitalization, %	30.6	44.9	1.7
	Average term of hospitalization, days	6.2	8.8	1.4
	Rate of lingering and complicated cases, %	3.9	13.8	3.5
10	The same index in in-patients, %	9.8	26.2	2.7
	Number of temporary disablement cases/100 persons/month	4.1	7.0	1.7
	Number of temporary disablement days/100 persons/month	26.5	57.6	2.2

**THE DYNAMICS OF ARD AND INFLUENZA RATE IN GROUPS OF OBSERVATION  
BY MONTHS FROM THE BEGINNING OF INVESTIGATION**

	Indices	Groups	1st month	2nd month	3rd month	4th month
20	Sickness rate/100 persons/month	THYMOGEN Control	9.6 28.6	11.3 33.4	9.4 28.7	11.0 30.6
	Need in hospitalization, %	THYMOGEN Control	27.0 41.2	27.1 50.8	28.3 48.8	28.3 39.0
	Average term of hospitalization, days	THYMOGEN Control	6.2 9.3	6.2 8.4	6.2 10.2	7.0 11.0
25	Number of temporary disablement cases/100 persons/month	THYMOGEN Control	3.8 6.9	7.3 10.2	3.4 7.5	3.6 5.6
	Number of temporary disablement days/100 persons/month	THYMOGEN Control	24.0 47.9	48.6 82.0	14.3 51.4	26.3 42.8

**EXAMPLE 9**

In separate studies, a total of 21 AIDS infected individuals have been studied, including full-blown syndrome, prodromal, and pre-AIDS afflicted individuals who were treated with Thympentin. Thympentin and TPI are thymic gland peptide extracts previously well characterized. Comparative studies

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between TPI and Thymogen reveal that Thymogen is a far more effective cell mediator, restoring normal immunologic indices, including T-cell functional activity and T4/T8 ratios. Method of Administration:

- 5 Sterile saline containing the sodium salt of the medication is administered either IM, intralymphatically, or intranasally each day for 5 - 10 days consecutively every 30 days.

- 10 Immunosuppressed individuals who have sustained radiation injuries were treated with Thymogen with excellent restoration of immunological indices and models for acquired immune deficiency syndrome. Thymogen may thus benefit AIDS infected individuals by reducing the need to use other medications with  
15 toxic side effects, and sustain and or support the individuals by reducing the needs to use other medications with toxic side effects, and sustain and or support the individuals immune indices resulting in a reduction of opportunistic infections.

20

EXAMPLE 10

- A total of 159 patients were treated with pyoderma, including furunculitis, cellulitis, and folliculitis, with a control group consisting of 25 patients who were not treated with thymogen.
- 25 Medications were administered either IM or intranasally for 5 consecutive days. Immunological indices were normalized with disappearance of skin manifestations and relapses were prevented after treatment with Thymogen. Clinical improvement  
30 correlated with immunological indices correction. Administration IM, intranasally, or topically as a sterile saline solution of medication for a period of 5 to 10 days at a concentration of 1  $\mu$ g/kg body weight.

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EXAMPLE 11

A number of patients within the group of 159 patients afflicted with furunculitis, pyoderma, cellulitis, and folliculitis were afflicted with acne vulgaris and acne. The immunological indices corrected and normalized rapidly within the group therapy. The clinical outcome correlated with the correction of immunological indices, and relapses were controlled.

10

EXAMPLE 12

A total of 30 patients were treated with psoriasis and 30 patients were used as controls and were untreated with thymogen. All patients had at a least 5 year history of no unsuccessful treatment. The administration of 100  $\mu$ g IM or intranasally for a period of 10 days resulted in the improvement in 7% of the patients, significant improvement in 60% of patients, and total recovery in 33% of the patients.

EXAMPLE 13

A total of 46 female patients with the various disorders (pelvic inflammatory diseases, cervicitis, vaginitis and various tubo-ovarian and adnexal abscesses) were treated and 50 patients were used as controls. Thymogen was applied IM, intranasally at 100  $\mu$ g 5 consecutive days or 50  $\mu$ g intralymphatically for 5 consecutive days in conjunction with conventional therapy. The clinical effect of Thymogen expressed the arresting of pain syndrome, the control of body temperature, e.g. reduction of fever, the decrease of duration of conventional

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treatment. The normalization of immune status correlated with clinical improvements.

#### EXAMPLE 14

Patients treated with Thymogen either topically,  
5 IM, or intranasally experienced marked reduction of  
recurrence of herpetic lesions, with substantial  
reduction in the period between outbreaks. In one  
study, individuals who experienced 7-10 outbreaks per  
year experienced less than one outbreak per year  
10 after treatment with Thymogen in combination with  
interferon.

#### EXAMPLE 15

A total of 37 patients with Herpes Zoster were  
treated with Thymogen in combination with  
15 conventional interferon treatment and 25 control  
patients with interferon alone. Administration  
single daily IM or intranasal 100  $\mu$ g during a period  
of 10 days resulted in accelerated regression of foci  
of herpetic infection. There was noted prevention of  
20 relapses, and healing occurred on the average 40%  
earlier than control groups. Immunological indices  
correlated with clinical outcome.

#### EXAMPLE 16

Patients were treated for gingival disease by  
25 subcutaneous administration of thymogen in the area  
of the gingiva. The treatment resulted in the  
arresting of gingival disease. Approximately 80  
patients were studied with disease and treated and an  
equal number were treated conventionally without  
30 Thymogen for control purposes. Administration of 100  
ug Im, Subcutaneously, or by electrophoresis (whereby

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a small voltage charge to the gums results in a rapid transfer of medication through the gum epithelium) resulted in the arresting of bleeding, more rapid restoration of inflammatory processes, and the  
5 decrease of purulent discharge. The treatment resulted in fewer recurrences and prolongation of normal gums. It was also noted that normalization of immunologic indices was achieved with normal coagulation.

10

EXAMPLE 17

The treatment with toothpaste containing Thymogen will result in a reduction of dental caries.

EXAMPLE 18

A total of 46 patients with periapical granulomas  
15 and 28 patients with the same disease not treated with Thymogen were used for controls. Instillation of 100  $\mu$ g of Thymogen into the foramen at the base of the tooth, or in the composition of the filling paste during 3 days resulted in the accelerated arrestation  
20 of the inflammatory process, reduction in pain, and increased stability of the underlying dental structures as evidenced by x-ray studies.

EXAMPLE 19

The use of dental toothpaste containing Thymogen  
25 will result in the reduction of gingival disease and reduction in dental caries.



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EXAMPLE 20

The use of Thymogen 100  $\mu$ g IM, intranasally, or intralymphatically controls the advance of lymphangitis.

5

EXAMPLE 21

A total of 186 patients with acute respiratory disease, including upper airway diseases, such as colds, were treated with Thymogen and 87 patients who were not treated with Thymogen were used as controls.

10 Administration IM or intranasally 100  $\mu$ g 3 - 7 days resulted in a milder course of the viral infection. There was noted a decrease in the specific signs of upper respiratory infections such rhinorrhea, sore throat, fever, muscle aches, headaches, and ear pain.

15 Secondary infectious complications were diminished, and the duration of the treatment was also diminished.

EXAMPLE 22

A total of 51 patients were treated with Thymogen

20 with 24 patient controls, administration IM, intranasally, and installation into sinuses with 1  $\mu$ g/kg dose during a period of 3 - 10 days resulted in normalization of nasal breathing, the disappearance of nasal mucous swelling, the arresting of exudates

25 from affect sinuses, and improved general condition and immune status. The decrease of treatment duration up to 1.7 times compared to controls.

EXAMPLE 23

Thymogen IM or intranasal accompanying

30 conventional therapy (antibiotics) results in

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accelerated healing of chronic and acute ear infections.

EXAMPLE 24

5 A total of 41 patients with various eye problems as described and 36 patients in control studies were treated by conventional methods with the first group receiving Thymogen in addition to the conventional treatment. Administration of Thymogen intra ocularly at 18  $\mu$ g for 5 consecutive days, or as installation  
10 into conjunctival cavity as drops bid for 5 days resulted in more rapid arresting of the inflammatory process and the increase in visual acuity, and the decrease of duration of treatment.

EXAMPLE 25

15 A total of 156 patients treated with Thymogen and 82 patients in the control study, were administered, medication IM or intranasally 100  $\mu$ g 5 - 10 days resulting in accelerated reduction in symptom complexes including joint pain, muscle aches, fevers,  
20 chills, and upper respiratory symptoms.

EXAMPLE 26

A total of 263 patients and 18 control patients sustained exposure to radiation injury. Thymogen was administered IM and/or intranasally 100  $\mu$ g for 10  
25 days. A repeated course may be prescribed on the basis of immunological indices, and averages every 4 to 6 months. The results of the treatment are restoration of normal or near normal immune indices with functional activity in the majority of all cases  
30 studied. There was an arresting of esthetic syndrome, and an arresting of the somatic

-25-

pathological exacerbations and reduction of opportunistic infections.

EXAMPLE 27

Thymogen administration IM or intranasally results  
5 in the improved immune parameters, functionals  
activity of lymphocytes and neutrophils, and  
reduction of post-operative complications and  
infections associated with bone-marrow compromise,  
such as, that caused from transplant or radiation  
10 exposure.

EXAMPLE 28

A total of 29 patients afflicted with various  
allergies as described and 17 patients in the control  
group were treated with Thymogen in dose 1  $\mu\text{g/kg}$  IM  
15 or intranasally for 5 - 7 days resulted in  
disappearance of allergic reactions.

EXAMPLE 29

A total of 76 patients with 72 patients in control  
exposed to massive hemotransfusions during post-  
20 operative period were treated with Thymogen.  
Thymogen was administered starting from 4-6 day of  
post-operative period single daily IM or intranasally  
in doses 100  $\mu\text{g}$  for 5 days. None of studies patients  
showed clinical manifestation of alloblood rejection  
25 while in 17% of control patients the adverse  
hemotranfusal reactions were observed.

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EXAMPLE 30

Thymogen was applied in 76 patients treated with antibiotics for various indications who had unfavorable allergological history. Control group comprised 43 patients. Thymogen was administered IM or intranasally single daily at 100  $\mu$ g for 5-10 days. In the majority of case the use of Thymogen prevented the arising of allergic reactions or promoted the less severe course of them while in the control group in 70% of patients the pronounced signs of drug intolerance was marked.

EXAMPLE 31

Thymogen was administered to 17 patients subjected to skin grafting. The control group comprised 27 patients. Thymogen was administered IM or intranasally single daily at 50-100  $\mu$ g for 5 days. In all the patients the use of Thymogen prevented the arising of infections complications and graft rejection. In control group the manifestations of rejection were determined in 8 patients.

EXAMPLE 32

Thymogen was administered to 52 patients suffered from chronic skin diseases caused by antibiotic-resistant staphylococci. 42 patients with the same pathology but not treated with the immunomodulator were the control group. Thymogen was administered IM to 27 patients single daily at 100  $\mu$ g for 5 days and intranasally to 25 patients in the same daily and total dose. The differences between these two methods of application were not noticed. In all the patients with signs of secondary T-immunodeficiency the staphylococci antibiotic-sensitivity to one, few or

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all antibiotics has been increased sharply (more than 100-fold) what permitted further to choose for each patient the antibiotic with exclusively high activity against pathogen. As a whole, within the examined  
5 group of patients the reliable decrease of MIC of all studied antibiotics has been marked. The proposed treatment regiment permitted to obtain the complete recovery in 27 patients, significant improvement - in 8 patients and moderate improvement - in 1 patient.

10

EXAMPLE 33

Thymogen was used in 37 patients with wounds of various origin, type and localization. the control group comprised 24 patients. Thymogen was  
15 administered IM or topically single daily at 100 ug for 10 days. The use of Thymogen speeded up (when compared to the control group) significantly wound healing, reduced therapy duration and prevented the development of infectious complications.

EXAMPLE 34

20 Administration of Thymogen either intranasally or IM accelerates wound healing, resulting in statistically fewer infections and reduced eschar.

EXAMPLE 35

Thymogen was applied to 44 patients with bone  
25 fractures various origin. type and localization. The control group comprised 28 patients. Thymogen was administered intramuscularly or intranasally single daily at 100 up for 10 days. The use of Thymogen accelerated essentially (in comparison with  
30 the control group) the consolidation of fractures, prevented the development of infectious

-28-

complications, reduced pain syndrome and treatment duration.

EXAMPLE 36

Thymogen was prescribed to 176 patients with  
5 chronic osteomyelitis of various ethiology and  
localization. The control group comprised 88  
patients. Thymogen was administered IM or  
intranasally single daily at 100 ug for 10 days. The  
use of thymogen rendered the pronounced positive  
10 influence on clinical course what expressed in  
significant decrease of intoxication syndrome and  
pain syndrome, disappearance of purulent inflammatory  
manifestations, speeding up of wound healing,  
reduction of destruction areas, prevention of  
15 relapses.

EXAMPLE 37

A total of 23 patients with cutaneous burns were  
treated with Thymogen either IM or intranasally with  
14 patients for control treated conventionally.  
20 Accelerated wound healing, diminished frequency of  
infections, and less eschar was noted in those  
individuals treated with Thymogen.

EXAMPLE 38

A total of 17 patients with frostbite to the  
25 extremities where treated with Thymogen either IM or  
intranasally with 11 patient controls. The rapid  
healing and restoration of tissue integrity was  
observed.

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EXAMPLE 39

Thymogen administration either IM or intranasally results in less deformity and scarring evidenced by experience in healing fractures, burns, military accidents, and other injuries to the extremities.

EXAMPLE 40

Experimental data supports the finding that Thymogen administered IM, intranasally, or ocular installation results in restoration and regeneration of corneal epithelium with fewer infections and complications related to eschar.

EXAMPLE 41

A total of 246 patients with various forms of cancer, and 158 controls after radiation and chemotherapy, where Thymogen was administered in single 100 ug daily dose for 10 days experienced normalization of immunological indices, the prevention of post-operative infections, the prevention of upper respiratory infections, and prevention of exacerbations of various secondary complications such as gastritis, cholecystitis, etc. If it was determined necessary based on immunological indices, the treatment regimen was repeated in 4-6 months.

EXAMPLE 42

Patients treated with Thymogen simultaneously during the administration of chemotherapy experienced fewer complications and side effects related to chemotherapy including diminished frequency and

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intensity of ulcerative lesions, nausea, and other related problems of chemotherapy administration.

EXAMPLE 43

Experimental models support the fact that  
5 administration of Thymogen prophylactically results in diminished frequency of spontaneous tumorigenesis.

EXAMPLE 44

Thymogen was applied to 268 persons in combination with the anti-flu vaccination. The control group  
10 comprised 197 persons. the vaccination was delivered by air pressure. The Thymogen dose was 50 ug delivered in a single dose for 3 consecutive days. After Thymogen use, it was observed the significant decrease of sickness rate for a period of 12 months  
15 compared to controls who received flu-vaccination without Thymogen. In the event of flu, the course of the infection was noted to be less severe and the recovery more rapid when compared to controls.

EXAMPLE 45

20 Thymogen was applied in 97 pregnant women with Toxemia of first and second half of pregnancy. The control group comprised 54 patients. Thymogen was administered IM and intranasally at 100 ug daily for 5 - 10 days. Under the influence of Thymogen, it was  
25 observed that the BP normalized, and peripheral edema was reduced with normalization of the blood chemistry profile, and the restoration of initially altered immunologic indices.



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EXAMPLE 46

Thymogen was administered to 34 pregnant women with 27 pregnant women for control. The route of administration is IM or intranasally 100 ug daily for 5 - 10 days. Signs of clinical improvement were resolution of weakness, dizziness, and increased appetite, and the normalization of the immunological and hematological indices. It was also noted that there was a decrease in fetal hypoxia.

10

EXAMPLE 47

A total of 19 post-term women and 48 women post-term in the control study were treated. The administration of 100 ug of Thymogen IM or intranasally over 3 - 5 days resulted in the effacement of the cervix with thinning at the cervix and the decent of the fetus, with subsequent spontaneous normal delivery.

EXAMPLE 48

A total of 27 patients with pyelonephritis and 19 control patients with pyelonephritis were treated with the administration of Thymogen single daily dose of 100 ug for 5 - 10 consecutive days in combination with conventional therapy resulted in reduction of fever, the normalization of urina analysis, and the improvement and resolution of the infection. The normal course of delivery in those women treated with Thymogen was without complications.

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EXAMPLE 49

A total of 45 patients with leprosy (Hansen's disease) and 27 infected individuals were treated. Thymogen was administered IM or intranasally in  
5 single daily doses of 100 ug for 5 days consecutively in additional to conventional therapy. The patients studied had previous documented resistance to treatment by conventional methods. Thymogen  
10 administration resulted in resolution of the lesions and prevented relapses, and promoted more rapid healing of specific ulcers. The immunologic indices were normalized.

EXAMPLE 50

Thymogen was administered to 84 young sportsmen.  
15 The control group consisted of 44 persons. The Thymogen was administered intranasally single dose 1 ug/kg during 3 days. The use of Thymogen resulted in the reduction of upper respiratory infections and rates of illness 4 fold. In the event of infection,  
20 it was noted that the infections was less severe without complications, and the clinical improvement was accompanied by the normalization of immunological indices.

EXAMPLE 51

25 A total of 33 patients were studied with patients who had relapsing forms of tropical malaria, moderate to severe, and severe cases with 21 patients in the control group. Thymogen was administered at 100 ug single daily doses IM or intranasally for 5 - 10  
30 days. the results of such treatment was reduction of hepatolinal syndrome, the normalization of

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hematological and immunological indices, reduction of fever, and prevention of relapses.

EXAMPLE 52

Thymogen was applied in 27 persons with the goal  
5 to increase the resistance to excessive solar radiation, in the conditions of hot marine climates. The control group comprised 24 persons. The administration was intranasally 100 ug for 3 days. The use of Thymogen prevented the occurrence of upper  
10 respiratory infections in the treated group relative to the control group. There was also noted suppression of their immunologic indices.

EXAMPLE 53

Thymogen was applied in 21 patients with  
15 hemorrhagic Dengue Fever, and 28 patients served as controls. Thymogen was administered IM single daily doses of 100 ug for 5 consecutive days in conjunction with conventional therapy. The results of treatment were reduction in fever, reduction of toxic symptoms,  
20 significant decrease in hepato-lineal syndrome. It was also noted that the muscular and bone pain experienced typically was reduced, and the immunological indices were normalized.

EXAMPLE 54

25 A total of 48 patients infected and 34 infected controls were examined and treated with administration of Thymogen 100 ug IM or intranasally for 5 - 10 days resulting in normalization of fever, the reduction of toxic symptoms, and the resolution  
30 of icterus (jaundice). The hematological and immunological indices were normalized.

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EXAMPLE 55

A total of 36 patients infected and 24 patients infected were controls. Administration of Thymogen in 100 ug IM or intranasally for 5 - 10 days resulted  
5 in the reduction of fever, more rapid reduction of toxic symptoms, and the restoration of immunologic indices.

EXAMPLE 56

A total of 37 patients infected with pulmonary TB  
10 and 26 patients infected as controls were studied and treated. Thymogen was administered at 50 to 100 ug every other day during 5 doses total in combination with convention therapy. The results of the  
15 treatment 2 months after the course of Thymogen revealed the disappearance of toxic symptoms, the reabsorption of infiltrates, and resolution of pulmonary cavities. The disappearance of TB bacilli was noted in the sputum. The restoration of initially decreased immune indices was also noted.

EXAMPLE 57

20 A total of 37 patients, children and adults, with bronchial asthma and 28 similar patients as controls were studied. Thymogen was administered IM single daily doses 1 ug/kg for 5 - 10 days resulting in less  
25 severe clinical symptoms. The significant reduction in bronchial obstruction and laryngotracheitis was noted. The normalization of fever, and the reduction in duration of treatment was noted. In some of the patients it was possible to avoid steroids in the  
30 conventional commitment treatment course. In the following year observation there was noted a decrease in the incidence of bronchial asthma 4.2 fold. In

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more than half of the patients the disappearance of drug and food allergy manifestations was noted.

EXAMPLE 58

A total 125 patients with 53 patients for control  
5 infected with Shigella dysentery were examined.  
Thymogen was administered IM single doses of 100 ug  
for 10 consecutive days with resultant normalization  
of fever, the reduction of toxemia, and the  
normalization gastrointestinal disorders and  
10 symptoms. Bacterial shedding in the GI track was  
observed to cease, and the immunological indices were  
normalized.

EXAMPLE 59

A total of 12 patients who had been thymectomized  
15 were treated with Thymogen. Prior to therapy these  
individuals had experienced frequent serious  
infections including upper respiratory infections.  
Thymogen was administered in a single dose 100 ug  
daily for 10 days and repeated every 4 - 6 months.  
20 The normalization of immunologic indices was  
observed, and there was reduction of infectious  
disorders including cutaneous infections and other  
chronic exacerbations.

EXAMPLE 60

25 A total of 39 patients were studied with 27  
patient controls. Thymogen was administered IM or  
intranasally at 100 up for 5 - 10 days to the study  
group of patients with the resulting reduction of  
fever, decrease in toxic symptoms, the reduction of  
30 musculoskeletal pain, and the reduction or

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disappearance of jaundice. Immunological indices were normalized.

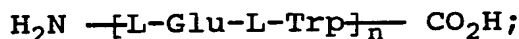
EXAMPLE 61

5 The use of Thymogen as an ingredient or applicant with cosmetics provides for a less allergenic cosmetic with fewer allergic reactions.

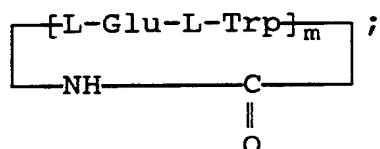
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WHAT IS CLAIMED IS:

1. A method for preventing and treating opportunistic infections in an immunodepressed or immunodeficient subject, comprising the step of  
 5 administering to said subject an effective immunopotentiating amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula



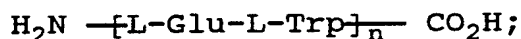
- 10 cyclic polymers thereof of the formula:



15

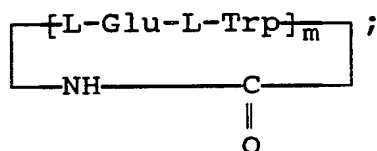
and their pharmaceutically acceptable salts thereof, wherein n and m are independently  $\geq 2$ .

2. A method for the treatment of an immunodeficient, immunodepressed or hyperactive  
 20 immune state in a subject comprising the step of administering to said subject an effective amount of a dipeptide selected from the group consisting of L-Glu-Trp the cyclic monomer thereof, polymers thereof of the formula



25

cyclic polymers thereof of the formula:



30

and their pharmaceutically acceptable salts thereof, sufficient to alleviate said immunodeficient, immunodepressed or hyperactive immune state, wherein  
 35 n and m are independently  $\geq 2$ .

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3. A method according to Claim 1 or 2 wherein said infection comprises acquired immune deficiency syndrome.

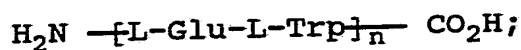
4. A method according to Claim 1 wherein said  
5 infection comprises skin disease.

5. A method according to Claim 4 wherein said skin disease is selected from the group consisting of pyoderma, furunculitis, cellulitis.

6. A method according to Claim 4 wherein said  
10 skin disease comprises eczema.

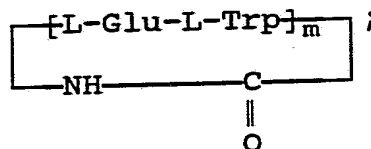
7. A method according to Claim 4 wherein said skin disease comprises acne, including acne vulgaris.

8. A method for treating psoriasis comprising the step of administering to a subject an effective  
15 amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula



cyclic polymers thereof of the formula:

20



25 and their pharmaceutically acceptable salts thereof, sufficient to alleviate said psoriasis, wherein n and m are independently  $\geq 2$ .

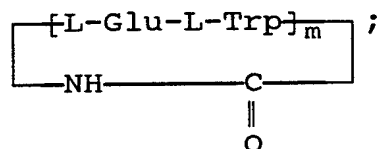
9. A method according to Claim 1 wherein said infection comprises a gynecological infection.



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10. A method according to Claim 9 wherein said infection is selected from the group comprising of pelvic inflammatory disease, cervicitis, vaginitis, and tubo-ovarian abscesses and adnexal abscesses.
- 5 11. A method according to Claim 1 wherein said infection comprises herpetic lesions.
12. A method according to Claim 11 wherein said infection comprises herpetic lesions of Type I or Type II categories.
- 10 13. A method according Claim 11 wherein said infection comprises herpetic lesions of Herpes Zoster.
14. A method according to Claim 1 wherein said infection comprises gingival disease.
- 15 15. A method according to Claim 1 wherein said infection comprises dental caries.
16. A method according to Claim 1 wherein said infection comprises periapical granulomas.
17. A method for prevention of dental caries or  
20 gingival disease in a subject comprising the step of administering to a subject an effective amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula

- 25 
$$\text{H}_2\text{N} - \{ \text{L-Glu-L-Trp} \}_n - \text{CO}_2\text{H};$$
  
cyclic polymers thereof of the formula:



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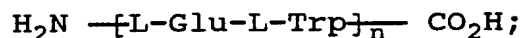
and their pharmaceutically acceptable salts thereof, sufficient to alleviate said dental caries or gingival disease, wherein n and m are independently  $\geq 2$ .

- 5 18. A method according to Claim 1 wherein said infection comprises infralymphatic infection.
19. A method according to Claim 18, where said infection comprises lymphangitis.
20. A method according to Claim 1 wherein said  
10 infection comprises acute respiratory disease.
21. A method according to Claim 1 wherein said infection comprises upper airway diseases, including the common cold.
22. A method according to Claim 1 wherein said  
15 infection comprises sinusitis and paranasitis.
23. A method according to Claim 1 wherein said infection comprises Otitis media and said dipeptide is co-administered with an antibiotic.
24. A method according to Claim 1 wherein said  
20 infection comprises conjunctivitis, uveitis, keratitis.
25. A method according to Claim 1 wherein said infection comprises influenza.
26. A method according to Claim 25 wherein said  
25 infection comprises influenza of category Type A or B, or any of their variants.

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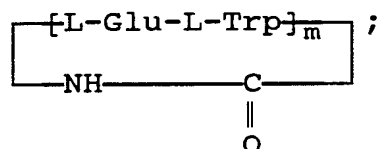
27. A method according to Claim 2 wherein said state is caused by exposure to radiation.

28. A method for bone marrow restoration comprising the step of administering to a subject an effective amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula



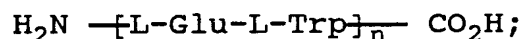
cyclic polymers thereof of the formula:

10



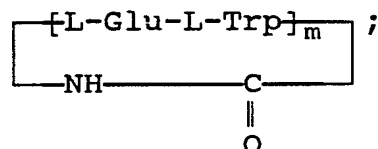
15 and their pharmaceutically acceptable salts thereof, sufficient to accelerate bone marrow restoration, wherein n and m are independently  $\geq 2$ .

29. A method for treatment of allergic reactions comprising the step of administering to a subject an effective amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula



cyclic polymers thereof of the formula:

25



30 and their pharmaceutically acceptable salts thereof, sufficient to alleviate said allergic reactions, wherein n and m are independently  $\geq 2$ .

30. A method for enhancing sensitivity of bacteria to antibiotics and for reducing the side effects to antibiotics comprising the step of

35

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administering to a subject an effective amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula

5 
$$\text{H}_2\text{N} - \{ \text{L-Glu-L-Trp} \}_n - \text{CO}_2\text{H};$$
  
cyclic polymers thereof of the formula:



and their pharmaceutically acceptable salts thereof, sufficient to enhance said sensitivity and to reduce said side effects, wherein n and m are independently  
15  $\geq 2$ .

31. A method for enhancing the healing process comprising the step of administering to a subject an effective amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer  
20 thereof, polymers thereof of the formula

$$\text{H}_2\text{N} - \{ \text{L-Glu-L-Trp} \}_n - \text{CO}_2\text{H};$$
  
cyclic polymers thereof of the formula:



and their pharmaceutically acceptable salts thereof, sufficient to enhance said process, wherein n and m  
30 are independently  $\geq 2$ .

32. A method for reducing scarring during healing comprising the step of administering to a subject an effective amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer  
35 thereof, polymers thereof of the formula

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$\text{H}_2\text{N} - \{\text{L-Glu-L-Trp}\}_n - \text{CO}_2\text{H};$   
 cyclic polymers thereof of the formula:



and their pharmaceutically acceptable salts thereof,  
 sufficient to reduce said scarring, wherein n and m  
 10 are independently  $\geq 2$ .

33. A method according to Claim 31 wherein said  
 treatment assists in the healing of local skin trauma  
 of skin grafts.

34. A method according to Claim 31 wherein said  
 15 treatment assists in healing of bone fractures.

35. A method according to Claim 31 wherein said  
 treatment improves the course of treatment for  
 osteomyelitis.

36. A method according to Claim 31 wherein said  
 20 treatment assists in the healing of burns and other  
 wounds.

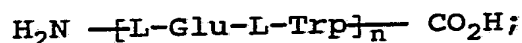
37. A method according to Claim 31 wherein said  
 treatment assists in the healing of frost bite.

38. A method according to Claim 31 wherein said  
 25 treatment promotes corneal regeneration and  
 restoration.

39. A method for prevention of spontaneous tumors  
 comprising the step of administering to a subject an  
 effective amount of a dipeptide selected from the

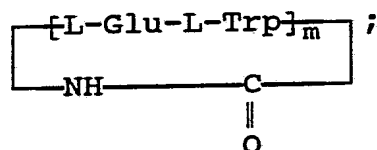
-44-

group consisting of L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula



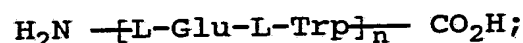
cyclic polymers thereof of the formula:

5



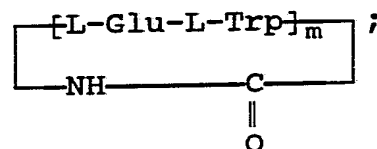
10 and their pharmaceutically acceptable salts thereof, sufficient to prevent spontaneous generation of said tumors, wherein n and m are independently  $\geq 2$ .

40. A method for enhancing the therapeutic effect of chemotherapy comprising the step of administering  
15 to a subject an effective amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula



20 cyclic polymers thereof of the formula:

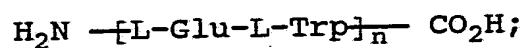
25



and their pharmaceutically acceptable salts thereof, sufficient to enhance said therapeutic effect, wherein n and m are independently  $\geq 2$ .

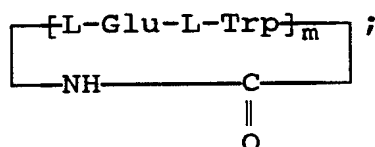
41. A method for enhancing the effect of  
30 vaccinations to a disease comprising the step of administering to a subject an effective amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula

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cyclic polymers thereof of the formula:

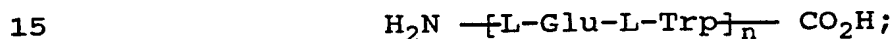
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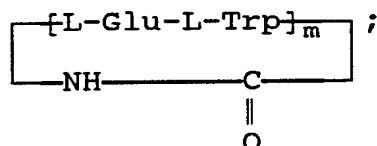
and their pharmaceutically acceptable salts thereof, sufficient to enhance said effect, wherein n and m are independently  $\geq 2$ .

42. A method for treating ailments associated with pregnancy comprising the step of administering to a subject an effective amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula



15

cyclic polymers thereof of the formula:



20

and their pharmaceutically acceptable salts thereof, sufficient to alleviate said ailment, wherein n and m are independently  $\geq 2$ .

25 43. A method according to Claim 42 for the treatment of toxemia of pregnancy.

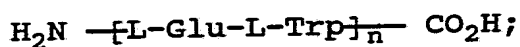
44. A method according to Claim 42 for the treatment of anemia of pregnancy.

45. A method according to Claim 42 to induce  
30 delivery in post-term pregnancies.

46. A method according to Claim 42 for the treatment of pyelonephritis during pregnancy in conjunction with conventional antibiotic treatment.

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47. A method for treatment of Hansen's disease comprising the step of administering to a subject an effective amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula



cyclic polymers thereof of the formula:



- and their pharmaceutically acceptable salts thereof, sufficient to alleviate said disease, wherein n and m are independently  $\geq 2$ .

48. A method according to Claim 2 wherein said immunodepression is related to stress induction of various types.

49. A method according to Claim 2 wherein said immunodepressed state is chronic fatigue syndrom.

50. A method according to Claim 1 wherein said infection comprises malaria.

51. A method according to claim 2 wherein said immunodepression is related to excess solar exposure.

- 25 52. A method according to Claim 1 for the treatment of hemorrhagic dengue fever.

53. A method according to Claim 1 for the treatment of Hepatitis A and B.



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54. A method according to Claim 1 for the treatment of typhus of the para A and B category.

55. A method according to Claim 1 for the treatment of tuberculosis of the lung.

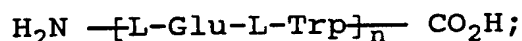
5 56. A method according to Claim 1 for the treatment of bronchial asthma.

57. A method according to Claim 1 for the treatment of shigella infected individuals with dysentery.

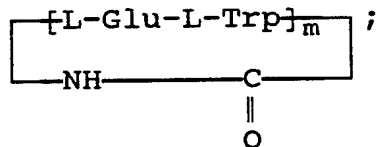
10 58. A method according to Claim 2 for the treatment of individuals who have been thymectomized.

59. A method according to Claim 1 for the treatment of yersenia, pseudo-tuberculosis.

60. A method for enhancing blood insulin  
15 comprising the step of administering to a subject an effective amount of a dipeptide selected from the group consisting of L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula



20 cyclic polymers thereof of the formula:



25

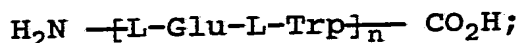
and their pharmaceutically acceptable salts thereof, sufficient to enhance said blood insulin, wherein n and m are independently  $\geq 2$ .

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61. A method according to Claim 2 wherein said treatment assists in recovery subsequent to anticancer radiation therapy.

62. A method according to Claim 1, 2, 28, 29, 30, 31, 39, 40, 42 or 59 wherein said subject is an animal.

63. A pharmaceutical preparation comprising a therapeutically effective amount to lower a hyperactive immune state of the dipeptide L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula

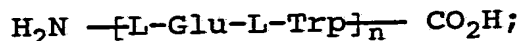


cyclic polymers thereof of the formula:



and/or their pharmaceutically acceptable salts thereof in a pharmaceutically acceptable vehicle, wherein n and m are independently  $\geq 2$ .

64. A pharmaceutical preparation for the restoration of normal immunological indices comprising a therapeutically effective amount to restore normal immunological indices in a subject of the dipeptide L-Glu-L-Trp the cyclic monomer thereof, polymers thereof of the formula



cyclic polymers thereof of the formula:



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and/or their pharmaceutically acceptable salts thereof in a pharmaceutically acceptable vehicle, wherein n and m are independently  $\geq 2$ .

65. A pharmaceutical preparation according to Claims 63 or 64 in the form of an injectable solution containing .001 to .01% by weight of said dipeptide.

66. A pharmaceutical preparation according to Claim 63 or 64 in the form of tablets, suppositories, capsules, eye films, inhalant, mucosal spray, toothpaste, ointments, and/or water soluble based creams.

67. A pharmaceutical preparation according to Claim 66 wherein said tablets, suppositories, capsules, eye films, inhalants, mucosal sprays, toothpaste, ointments, and/or water soluble based creams contain at least 0.01 mg of said dipeptide per unit preparation.

68. A substantially pure compound selected from the group consisting of the cyclic monomer of the dipeptide L-Glu-L-Trp, polymers thereof of the formula

$$\text{H}_2\text{N} - \{ \text{L-Glu-L-Trp} \}_n - \text{CO}_2\text{H},$$
  
cyclic polymers thereof of the formula



and their pharmaceutically acceptable salts; wherein n and m are independently  $\geq 2$ .

## INTERNATIONAL SEARCH REPORT

International Application No.  
PCT/US92/09114**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(5) : A61K 37/00; CO7K 5/00

US CL : 514/19; 530/331

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/19; 530/331

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS, CAS, BIOSIS, MEDLINE

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	Medline Abstracts, Issued 1990, Rodionov et al., "The immunocorrective Therapy of Pyoderma Caused By Staphylococci Multiply Resistant To Antibiotics", Abstract No. 90224329, Vestn. Dermatol. Venerol., Volume 1, pages 42-45. See Abstract	1,4,5,30,62-64,66-68
Y	Medline Abstract, Issued September 1991, Iakovlev et al., "The Biochemical And Immunological Indices In The Rehabilitation Period Of The Victims Of The Accident On The Komsomolets Atomic Submarine", Abstract No. 92101433, Voen. Med. Zh, Volume 9, pages 28-33. See Abstract	1,61,63-65,68
Y	Bulletin: "Thymogen", published 1989 by Cytomed (Leningrad), pages 3-10.	1-68
Y	Medline Abstract, Issued 1991, Khmel'nitskii et al., "Morphofunctional Characteristics Of The Immunocompetent System In Hypotrophy And Its Correction By Thymogen, Abstract No. 92171772, Arkh. Patol. Volume 53(10), pages 24-27. See Abstract.	1,62

☒ Further documents are listed in the continuation of Box C.
 ☐ See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be part of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

14 January 1993

Date of mailing of the international search report

26 JAN 1993

Name and mailing address of the ISA/  
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## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US92/09114

## C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	Medline Abstract, issued September-October 1991, Grigoriants et al., "Immunocorrection In The Combined Treatment Of Patients With Osteomyelitis Developing Following Combined Injuries To The Maxillofacial And Craniocerebral Areas", Abstract No. 92188368, Stomatologia(Mosk), Volume 5, pages 53-54. See Abstract.	1,2,35,62-64,68
Y	Biological Abstract, Volume 92, Issued 1991, Rodionov et al., "Natural Killer Activity In Patients With Chronic Dermatoses", Abstract No. 100385, Vestn. Dermatol. Venerol. Volume 5, pages 4-6. See Abstract.	1,2,4-8,62-68
Y	Chemical Abstracts, Volume 116, No. 17, Issued 1990, Aliev et al. "Simulation of Thymus Dysfunction In Guinea Pigs By Using Immunomodulators", Abstract No. 171986U, Izv. Akad. Nauk. Az. Ssr, Ser. Biol. Nauk., Volume 1, pages 73-80. See Abstract.	1-3, 63-68
Y	Chemical Abstracts, Volume 116, No. 17, Issued 1991, Demidov et al., "Effects Of Thymus Preparations And Antituberculous Drugs On Immunological Reactivity And The Course Of Tuberculous Process In Experimental Animals", Abstract No. 165824Y, Probl. Tuberk. Volume 12, pages 52-54. See Abstract.	1, 55, 62, 64-68